CLPB gene
ClpB homolog, mitochondrial AAA ATPase chaperonin

Normal Function

The *CLPB* gene provides instructions for making a protein whose function is unknown. The CLPB protein is found in cells throughout the body but is most abundant in the brain. Based on its similarity to a protein in other organisms, researchers speculate that the CLPB protein helps unfold misfolded proteins so they can be refolded correctly. When misfolded, proteins cannot function properly and may be damaging to cells.

Health Conditions Related to Genetic Changes

**CLPB deficiency**

At least 20 *CLPB* gene mutations have been found to cause CLPB deficiency. This condition is characterized by neurological problems, including movement abnormalities and seizures; a shortage of white blood cells (neutropenia), which can increase the risk of infections; and clouding of the lenses of the eyes (cataracts). In addition, affected individuals have an increased amount of a molecule called 3-methylglutaconic acid in their urine, which does not appear to cause health problems. The severity of these features varies widely. Many of the *CLPB* gene mutations lead to an abnormally short CLPB protein that is likely broken down quickly. Other mutations may reduce CLPB's function. The severity of the condition is thought to be related to the amount of functional protein remaining: severe CLPB deficiency is likely caused by a complete absence of CLPB protein, while moderate and mild CLPB deficiency result when some functional CLPB protein is produced. Researchers are unsure how reduction or absence of this protein leads to the signs and symptoms of CLPB deficiency.
Chromosomal Location

Cytogenetic Location: 11q13.4, which is the long (q) arm of chromosome 11 at position 13.4

Molecular Location: base pairs 72,285,495 to 72,434,684 on chromosome 11 (Homo sapiens Updated Annotation Release 109.20190905, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- ANKCLB
- ankyrin-repeat containing bacterial clp fusion
- caseinolytic peptidase B protein homolog isoform 1
- caseinolytic peptidase B protein homolog isoform 2
- caseinolytic peptidase B protein homolog isoform 3
- caseinolytic peptidase B protein homolog isoform 4
- ClpB caseinolytic peptidase B homolog
- FLJ13152
- HSP78
- SKD3
- suppressor of potassium transport defect 3
- testicular secretory protein Li 11

Additional Information & Resources

Educational Resources

  https://www.ncbi.nlm.nih.gov/books/NBK21750/
Clinical Information from GeneReviews

• CLPB Deficiency
  https://www.ncbi.nlm.nih.gov/books/NBK396257

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28CLPB%5BTIAB%5D%29+OR+%28ClpB+homolog,+mitochondrial+AAA+ATPase+chaperonin%5BTIAB%5D%29+OR+%28%28ClpB+caseinolytic+peptidase+B+homolog%5BTIAB%5D%29+OR+%28HSP78%5BTIAB%5D%29+OR+%28SKD3%5BTIAB%5D%29+OR+%28ankyrin-repeat+containing+bacterial+clp+fusion%5BTIAB%5D%29+OR+%28caseinolytic+peptidase+B+protein+homolog+isoform+1%5BTIAB%5D%29+OR+%28caseinolytic+peptidase+B+protein+homolog+isoform+2%5BTIAB%5D%29+OR+%28caseinolytic+peptidase+B+protein+homolog+isoform+3%5BTIAB%5D%29+OR+%28caseinolytic+peptidase+B+protein+homolog+isoform+4%5BTIAB%5D%29+OR+%28suppressor+of+potassium+transport+defect+3%5BTIAB%5D%29+OR+%28testicular+secretory+protein+Li+11%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• CASEINOLYTIC PEPTIDASE B
  http://omim.org/entry/616254

Research Resources

• Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_CLPB.html

• ClinVar

• HGNC Gene Symbol Report

• Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:81570

• NCBI Gene

• UniProt
  https://www.uniprot.org/uniprot/Q9H078
Sources for This Summary

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